

**AMENDMENTS TO THE CLAIMS**

Claims 1-26 canceled.

27. (Previously presented) The method of claim 80, wherein the spray composition further comprises a flavoring agent in the amount of between 0.1 and 10 percent by weight of the total composition.

28. (Previously presented) The method of claim 27, wherein the polar solvent is present in an amount between 27 and 98 percent by weight of the total composition, the active compound is present in an amount between 0.005 and 55 percent by weight of the total composition, and the flavoring agent is present in an amount between 0.5 and 8 percent by weight of the total composition.

29. (Previously presented) The method of claim 80, wherein the polar solvent is present in an amount between 59 and 97 percent by weight of the total composition, the active compound is present in an amount between 0.01 and 40 percent by weight of the total composition, and the flavoring agent is present in an amount between 0.75 and 7.5 percent by weight of the total composition.

30. (Previously presented) The method of claim 80, wherein the polar solvent is selected from the group consisting of polyethylene glycols having a molecular weight between 400 and 1000, C<sub>2</sub> to C<sub>8</sub> mono- and poly-alcohols or C<sub>7</sub> to C<sub>18</sub> alcohols of linear or branched configuration.

31. (Previously presented) The method of claim 80, wherein the polar solvent comprises aqueous polyethylene glycol.

32. (Previously presented) The method of claim 80, wherein the polar solvent comprises aqueous ethanol.

33. (Canceled).

34. (Previously presented) The method of claim 80, wherein the flavoring agent comprises a synthetic or natural oil of peppermint, oil of spearmint, citrus oil, fruit flavors, sweeteners or mixture thereof.

Claims 35 – 53 canceled.

54. (Previously presented) The method of claim 81, further comprising a flavoring agent in the amount between 0.1 and 10 percent by weight of the total composition.

55. (Previously presented) The method of claim 54, wherein the non-polar solvent is present in an amount between 69 and 99 percent by weight of the total composition, the active compound is clozapine in an amount from between 0.5 and 30 percent by weight of the total composition, and the flavoring agent is present in an amount between 0.1 and 5 percent by weight of the total composition.

56. (Canceled).

57. (Previously presented) The method of claim 54, wherein the flavoring agent comprises a synthetic or natural oil of peppermint, oil of spearmint, citrus oil, fruit flavors, sweetener or a mixture thereof.

58. (Previously presented) The method of claim 81, wherein the solvent is selected from the group consisting of (C<sub>2</sub>-C<sub>24</sub>) fatty acid (C<sub>2</sub>-C<sub>6</sub>) esters, C<sub>7</sub>-C<sub>18</sub> hydrocarbons of linear or branched configuration, and C<sub>2</sub>-C<sub>6</sub> alkanoyl esters or triglycerides of a C<sub>2</sub>-C<sub>6</sub> carboxylic acid.

59. (Previously presented) The method of claim 81, wherein the solvent comprises one or more fatty acid esters.

Claims 60 – 79 canceled.

80. (Currently amended) A method for administering an effective amount of a pharmacologically active compound to a mammal to provide transmucosal absorption of a pharmacologically effective amount of the active compound through the oral mucosa of the mammal to the systemic circulatory system of the mammal, comprising:

spraying the oral mucosa of the mammal with a propellant free buccal spray composition, containing a pharmacologically active compound dissolved in a pharmacologically acceptable solvent, comprising in weight percent of the composition:

an active compound in an amount of between 0.001 and 60 percent selected from the group consisting of a ~~biologically active peptide, central nervous system active amine, sulfonyl urea, antibiotic, antifungal, sleep inducer, antiasthmatic, antiemetic, antiviral, histamine H<sub>2</sub> receptor antagonist, barbiturate, prostaglandin or bronchial dilator~~ clozapine, phenytoin or a pharmaceutically acceptable salt thereof; and

a polar solvent in an amount between 30 and 99.69 percent.

81. (Currently amended) A method for administering an effective amount of a pharmacologically active compound to a mammal to provide transmucosal absorption of a pharmacologically effective amount of the active compound through the oral mucosa of the mammal to the systemic circulatory system of the mammal, comprising:

spraying the oral mucosa of the mammal with a propellant free buccal spray composition, containing a pharmacologically active compound dissolved in a pharmacologically acceptable solvent, comprising in weight percent of the composition:

an active compound in an amount between 0.005 and 55 percent comprising selected from the group consisting of a biologically active peptide, central nervous system active amine, sulfonyl urea, antibiotic, antifungal, sleep inducer, antiasthmatic, antiemetic, antiviral, histamine H-2 receptor antagonist, barbiturate, prostaglandin or bronchial dilator clozapine, phenytoin or a pharmaceutically acceptable salt thereof; and

a non-polar solvent in an amount between 30 and 99.69 percent.

82. (Currently amended) A method for administering an effective amount of a pharmacologically active compound to a mammal to provide transmucosal absorption of a pharmacologically effective amount of the active compound through the oral mucosa of the mammal to the systemic circulatory system of the mammal, comprising:

spraying the oral mucosa of the mammal with a propellant free buccal spray composition, containing a pharmacologically active compound dissolved in a pharmacologically acceptable solvent, comprising in weight percent of the composition:

an active compound in an amount between 0.005 and 55 percent comprising ~~an antihistamine, alkaloid, hormone, a benzodiazepine or narcotic analgesic~~; and

a non-polar solvent in an amount between 30 and 99.69 percent.